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The importance of differentiating behavioural and psychological treatment effects from placebo in respiratory interventions.

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@ERSpublications: Well-designed non-pharmacological interventions can harness treatment responses seen in the placebo arm.

We read with interest the piece by Pattinson and Wanigaseraka [1], discussing the clinical implications of the recent trial by Currow et al. [2], which demonstrated parity between sertraline and placebo for improving breathlessness intensity. In particular, Pattinson and Wanigaseraka noted that differential response expectations can potentially confound outcomes between otherwise balanced intervention vs. placebo-controlled groups. Such expectations have clinical implications: the management of response expectations should be examined and potentially harnessed to improve treatment outcomes for patients with chronic breathlessness, as suggested by Similowski and Serresse [3].

In his excellent paper, Turner [4] reiterated that a placebo-controlled trial (PCT) compares a particular component of a treatment (often primarily pharmacological) while attempting to keep all other therapeutically-relevant components identical. Ideally, such an approach precludes bias and is the gold-standard to develop evidence-based medical treatments. However, chronic breathlessness (amongst other complex long-term conditions) is increasingly recognised as dysfunction across multiple interconnected systems. As such it requires holistic treatment that, alongside physiological factors, addresses social, psychological, neurocognitive and behavioural aspects of disease. Therefore, it must be acknowledged that although PCTs are designed to examine one specific treatment component, multiple components will impact patient outcomes – and these components may be present in both the treatment and placebo group.

This issue is particularly relevant in respiratory research, in which studies have consistently demonstrated very limited associations between objective physiological impairment and the subjective perceptions of breathlessness severity [5,6]. Thus, treatment benefits may impact patients through a diverse and complex interacting set of mechanisms (such as neurocognitive predispositions towards subjective symptom severity, as suggested by Ongaro and Kaptchuk [7]) which Pattinson and Wanigaseraka rightly note may not be balanced between PCT trial arms. Pertinently, we do not wish to devalue ‘traditional’ RCTs. However, evidence from such trials should be considered alongside other types of research methods, such as ‘pragmatic’ research that more

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3 closely reflects routine clinical care and ‘additively’ evaluates the incorporation of a new
4 treatment to a patients’ existing care, carefully monitoring all aspects of the new
5 treatment (e.g. interaction with healthcare, subsequent self-management) in order to
6 understand the likely mechanisms by which treatment benefits are conferred.
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9 Many medical professionals think of a ‘placebo’ as an inert pill but, as Turner noted, a
10 comparison with ‘placebo’ in a trial is merely a comparison of one group with another –
11 and the ‘placebo pill’ is merely a symbolic object involved in this process. This process
12 definition is in line with modern accounts of clinical placebos focussed on context,
13 meaning and embodiment. We also agree with Pattinson and Wanigaseraka’s point that
14 ‘placebo’ is often used disparagingly to refer to treatments that do not work; such a
15 position risks overlooking the benefits that are conferred independently to identified
16 physiological improvements. This is particularly relevant for non-pharmacological
17 treatments such as breathing retraining exercises, mindfulness-based treatments or
18 cognitive behavioural therapy, which are increasingly used as effective patient
19 treatments in respiratory disease but demonstrate no measurable effect on
20 physiological outcomes. By defining placebos as ‘inert’ substances we are in danger of
21 classifying such complementary treatments as ineffective, when in fact they may be
22 cost-effective adjunct treatments that offer tangible patient benefits.
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27 We welcome the call for ‘open-label’ placebo research. However, a considered and
28 cautious approach should be taken, insofar as such treatment may merely demonstrate
29 the presence of broad psychological and behavioural treatment effects without
30 determining which of these can be targeted (and therefore impacted by well-designed
31 non-pharmacological interventions). Moreover, as Currow[8] notes (supported by
32 modern contextual accounts of the placebo phenomenon[9]), although the results of
33 small-scale open-label placebo trials may reflect genuine therapeutic benefit, they may
34 also just be an artefact of the experimental situation – meaning it is unclear how such a
35 treatment reponse can be harnessed.
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38 These effects of open-label placebos are not caused by the ‘inert’ pill itself, but by the
39 construction and exploitation of a whole treatment process. Given that many existing
40 non-pharmacological interventions can be conceived of as already established whole
41 treatment processes (without the negative connotations of placebos) we advocate that
42 the best way to therapeutically employ beneficial treatment responses may be to
43 conduct careful research on these non-pharmacological interventions.
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47 **References**
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